Supporting Information

The Journal of Organic Chemistry

Synthesis and Biological Evaluation of Aromatic Analogues of Conduritol F, L-chiro-Inositol and Dihydroconduritol F Structurally Related to the Amaryllidaceae Anticancer Constituents

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General Methods. Unless otherwise noted all commercially obtained reagents were used without purification. THF was distilled from sodium-benzophenone ketyl prior to use. Dichloromethane and methanol were distilled from calcium hydride. Reactions were carried out under a nitrogen atmosphere in oven-dried glassware using standard syringe, cannula and septa techniques. Reactions were monitored by TLC (Silica Gel 60 F254, 250 μ m) and visualized with UV light and ceric ammonium molybdate solution. Flash chromatography was performed on silica gel (32-63 μ m, 60 Å pore size). Aryl bromides \mathbf{c}^1 and \mathbf{e} , and MOMCl⁴ were prepared as previously described.











































































































































































